In the Claims:

Please amend the claims as follows:

1. (Currently Amended) A compound of the Formula I or II

$$R^4$$
 NH_2
 N

in which

- R^1 is hydrogen, or branched and unbranched C_1 - C_6 -alkyl, it also being possible for one C atom of the alkyl radical to carry OR^{11} or a group R^5 , where R^{11} is hydrogen or C_1 - C_4 -alkyl, and
- is hydrogen, chlorine, bromine, iodine, fluorine, CF3 CF3, nitro, NHCOR²¹, NR²²R²³, OH, O-C₁-C₄-alkyl, O-C₁-C₄-alkylphenyl, NH₂, CN, a straight or branched C₁-C₆-alkyl, OR²¹ or phenyl, it also being possible for the phenyl rings to be substituted by at most two radicals R²⁴, and R²¹ and R²², independently of one another, are hydrogen or C₁-C₄-alkyl, and R²³ is hydrogen, C₁-C₄-alkyl or phenyl, and R²⁴ is OH, C₁, C₆-alkyl C₁-C₆-alkyl, O-C₁-C₄-alkyl, chlorine, bromine, iodine, fluorine, CF₃, nitro or NH₂, and
- x may be 0, 1 or 2 and
- R³ is $-D (F^1)_p (E)_q (F^2)_r$, $G -D (F^1)_p (E)_q (F^2)_r G$, where p, q and r may not simultaneously be 0, or is $-E (D)_u (F^2)_s (G)_v E (D)_u (F^2)_s (G)_v$, it also being

possible for the radical E to be substituted by one or two radicals A, and if v = 0, E is imidazole, pyrrole, pyridine, pyrimidine, piperazine, pyrrolidine or piperidine, or R^3 is $\underline{-O-(CH_2)_o-(CHR^{31})_m-(CH_2)_n-G}$,

$$-N$$
 R^{32} , $-N$
 R^{32} , $-N$
 R^{52}

or R³ is B, and

R³¹ is hydrogen, C₁-C₄-alkyl, OH or O-C₁-C₄-alkyl and

 R^{32} is hydrogen, $-(CH_2)_0-(CHR^{31})_m-(CH_2)_n-G$ or $-(CH_2)_p-G$,

m and o independently of each other are 0, 1, or 2 and

n may be 1, 2, 3 or 4 and

is hydrogen, chlorine, fluorine, bromine, iodine, branched and unbranched C₁-C₆-alkyl, OH, nitro, CF₃, CN, NR⁴¹R⁴², NH-CO-R⁴³ or O-C₁-C₄-alkyl, where R⁴¹ and R⁴², independently of one another, are hydrogen or C₁-C₄-alkyl and

 R^{43} is hydrogen, C_1 - C_4 -alkyl, C_1 - C_4 -alkylphenyl or phenyl, and

- D is S or Θ O, and
- E is phenyl, imidazole, pyrrole, thiophene, pyridine, pyrimidine, piperazine, pyrazine, furan, thiazole, isoxazole, pyrrolidine, piperidine or trihydroazepine, and
- F¹ is a chain of 1 to 8 carbon atoms, it also being possible for one carbon atom of the chain to carry an OH or O-C₁-C₄-alkyl group, and
- F² is a chain of 1 to 8 carbon atoms, it also being possible for one carbon atom of the chain to carry an OH or O-C₁-C₄-alkyl group, and
- p may be 0 or 1,

- q may be 0 or 1,
- r may be 0 or 1,
- s may be 0 or 1,
- u may be 0 or $\frac{1}{2}$,
- v may be 0 or 1, and
- G may be $NR^{51}R^{52}$ or

$$R^{52}$$
 R^{52}
 R^{52}

and

R⁵¹ is hydrogen or branched and unbranched C₁-C₆-alkyl or (CH₂)_t-K, and

R⁵² is hydrogen, branched and unbranched C₁-C₆-alkyl, <u>COCH</u>₃, <u>COCF</u>₃, phenyl,

in which

R⁵³ may be branched or unbranched O-C₁-C₆-alkyl, phenyl or branched or unbranched C₁-C₄-alkylphenyl, where in the case of R⁵² and R⁵³, independently of one another, one hydrogen of the C₁-C₆-alkyl radical may be substituted by one of the following radicals: OH, O-C₁-C₄-alkyl, cyclohexyl, cyclopentyl, tetrahydronaphthyl, cyclopropyl, cyclobutyl, cycloheptyl, naphthyl and phenyl, it also being possible for

the carbocycles of the radicals R^{52} and R^{53} , independently of one another, to carry one or two of the following radicals: branched or unbranched C_1 - C_6 -alkyl, branched or unbranched O- C_1 - C_4 -alkyl, OH, F, Cl, Br, $\frac{1}{L}$, CF₃, $\frac{1}{N}$ 0, NH₂, CN, COOH, COOC₁- $\frac{1}{C_4}$ -alkyl, $\frac{1}{C_4}$ -alkylarnino $\frac{1}{C_1}$ - $\frac{1}{C_4}$ -alkylarnino, CCl₃, $\frac{1}{C_4}$ -dialkylarnino $\frac{1}{C_1}$ - $\frac{1}{C_4}$ -alkylarnino, SO₂- $\frac{1}{C_4}$ -alkyl, SO₂phenyl, CONH₂, CONH- $\frac{1}{C_4}$ -alkyl, CONHphenyl, CONH- $\frac{1}{C_4}$ -alkylphenyl, NHSO₂- $\frac{1}{C_4}$ -alkyl, NHSO₂phenyl, S- $\frac{1}{C_4}$ -alkyl,

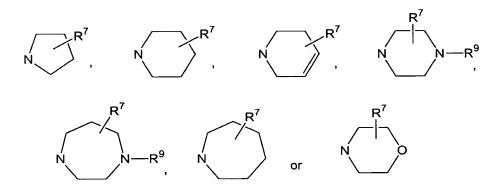
$$C_1$$
- C_4 -alkyl, C_0 - C_4 -alkylphenyl,

CHO, CH_2 -O- C_1 - C_4 -alkyl, $-CH_2$ O- C_1 - C_4 -alkylphenyl, $-CH_2$ OH, -SO- C_1 - C_4 -alkylphenyl, $-SO_2$ NH- C_1 - C_4 -alkyl, or two radicals form a bridge -O- $(CH_2)_{1,2}$ -O-,

B may be

and

- A may be hydrogen, chlorine, bromine, iodine, fluorine, CF₃, nitro, OH, O-C₁-C₄-alkyl, O-C₁-C₄-alkylphenyl, NH₂, branched and unbranched C₁-C₆-alkyl, CN or NH-CO-R³³, where R³³ is hydrogen, C₁-C₄-alkyl or phenyl, and
- t is 0, 1, 2, 3, or 4 and
- is phenyl, which may carry at most two radicals, is $NR^{k1}R^{k2}$ where R^{k1} and R^{k2} are as defined for R^{41} and R^{42} respectively, $NH-C_1-C_4$ -alkylphenyl, pyrrolidine, piperidine, 1,2,5,6-tetrahydropyridine, morpholine, trihydroazepine, piperazine, which may also be substituted by an alkyl radical C_1-C_6 -alkyl or homopiperazine, which may also be substituted by an alkyl radical C_1-C_6 -alkyl, and
- R⁵ may be hydrogen, C₁-C₆-alkyl, NR⁷R⁹ and



and

- R^7 is hydrogen, C_1 - C_6 -alkyl, C_1 - C_4 -alkylphenyl or phenyl, it also being possible for the rings to be substituted by up to two radicals R^{71} , and
- R⁷¹ is OH, C₁-C₆-alkyl, O-C₁-C₄-alkyl, chlorine, bromine, iodine, fluorine, CF₃, nitro or NH₂, and
- R⁸ is hydrogen, C_1 - C_6 -alkyl, phenyl or C_1 - C_4 -alkylphenyl C_1 - C_4 -alkylphenyl, it also being possible for the ring to be substituted by up to two radicals R⁸¹, and
- R⁸¹ is OH, C₁-C₆-alkyl, O-C₁-C₄-alkyl, chlorine, bromine, iodine, fluorine, CF₃, nitro, or NH₂, and
- is hydrogen, COCH₃, CO-O-C₁-C₄-alkyl, COCF₃, branched and unbranched C₁-C₆-alkyl, it being possible for one or two hydrogens of the C₁-C₆-alkyl radical to be substituted in each case by one of the following radicals: OH, O-C₁-C₄-alkyl or phenyl, and for the phenyl ring also to carry one or two of the following radicals: iodine, chlorine, bromine, fluorine, branched or unbranched C₁-C₆-alkyl, nitro, amino, C₁-C₄-alkylamino, C₁-C₄-dialkylamino, OH, O-C₁-C₄-alkyl, CN, CF₃ or SO₂-C₁-C₄-alkyl,

or a tautomeric form, a possible enantiomeric or disasteriomeric form, a prodrug or pharmacologically tolerated salt thereof.

- 2. (Currently Amended) A compound of the formula I or II as claimed in claim 1 in which
 - R^1 is hydrogen, branched and unbranched $C1 C_6$ -alkyl C_1 - C_6 -alkyl, it also being possible for one C atom of the alkyl radical to carry OR^{11} or a group R^5 , where
 - R¹¹ is hydrogen or C₁-C₄-alkyl, and
 - R^2 is hydrogen, chlorine, fluorine, bromine, iodine, branched and unbranched C_1 - C_6 alkyl, nitro, CF_3 , CN, $NR^{22}R^{23}$, NH-CO- R^{21} , or OR^{21} , where
 - R^{21} and R^{22} are, independently of one another, is hydrogen or C_1 - C_4 -alkyl, and

R23 is hydrogen or C₁-C₄ alkyl or phenyl, and

 R^3 is $-O-(CH_2)_o-(CHR^{31})_m-(CH_2)_n-G$, where

 R^{31} is hydrogen, C_1 - C_4 -alkyl, OH and or O- C_1 - C_4 -alkyl,

m and o are, independently of one another, 0, 1 or 2, and

n is 1, 2, 3 or 4, and

 R^4 is hydrogen, branched and unbranched C_1 - C_6 -alkyl, chlorine, bromine, fluorine, nitro, cyano, $NR^{41}R^{42}$, NH-CO- R^{43} , or OR^{41} , where

 R^{41} and R^{42} are, independently of one another, hydrogen or $C_1\text{-}C_4\text{-alkyl}$, and

 R^{43} is $C^{\frac{1}{2}}$ - C_4 -alkyl C_1 - C_4 -alkyl or phenyl, and

G is NR⁵¹R⁵² or one of the following radicals

$$R^{52}$$
 $N - R^{52}$ R^{52} R^{52} R^{52}

where

R⁵¹ is hydrogen and branched and unbranched C₁-C₆-alkyl, and

R⁵² hydrogen, branched and unbranched C₁-C₆-alkyl, phenyl,

$$\mathbb{R}^{53}$$
, or SO_2R^{53}

in which

is branched or unbranched O-C₁-C₆-alkyl, phenyl, branched or unbranched C₁-C₄-alkyl-phenyl, where one hydrogen in the C₁-C₆-alkyl radical in R⁵² and R⁵³ can, independently of one another, be substituted by one of the following radicals: OH, O-C₁-C₄-alkyl, cyclohexyl, cyclopentyl, tetrahydronaphthyl, cyclopropyl, cyclobutyl, cycloheptyl, naphthyl and phenyl, where the carbocycles of the R⁵² and R⁵³ radicals may also, independently of one another, carry one or two of the following radicals: branched or unbranched C₁-C₆-alkyl, branched or unbranched O-C₁-C₄-alkyl, OH, F, G₁ Cl, Br, ‡ I, CF₃, NO₂ NO₂, NH₂, CN, COOH, COOC₁-C₄-alkyl, C₁-C₄-alkylamino, CCl₃, C₁-C₄-dialkylamino, SO₂-C₁-C₄-alkyl, SO₂phenyl, CONH₂, CONH-C₁-C₄-alkyl, CONHphenyl, CONH-C₁-C₄-alkyl-phenyl, NHSO₂-C₁-C₄-alkyl, NHSO₂phenyl, S-C₁-C₄-alkyl,

$$C_{1}$$
- C_{4} -alkyl , C_{0} - C_{4} -alkylphenyl ,

CHO, CH_2 -O- C_1 - C_4 -alkyl, - CH_2 O- C_1 - C_4 -alkyl-phenyl, - CH_2 OH, -SO- C_1 - C_4 -alkyl-phenyl, SO₂NH₂, or -SO₂NH- C_1 - C_4 -alkyl

and two radicals form a bridge -O-(CH₂)_{1,2}-O-,

or the tautomeric form, possible enantiomeric and diastereomeric forms thereof, the prodrugs thereof, and pharmacologically physiologically tolerated salts thereof.

- 3. (Currently Amended) A compound of the formula I or II as claimed in claim 1 in which
 - R^1 is hydrogen, branched and unbranched C_1 - C_6 -alkyl, it also being possible for one C atom of the alkyl radical to carry OR^{11} or a group R^5 , where
 - R¹¹ is hydrogen or C₁-C₄-alkyl, and

 R^2 is hydrogen, chlorine, fluorine, bromine, iodine, branched and unbranched C_1 - C_6 -alkyl, nitro, CF_3 , CN, $NR^{22}R^{23}$, NH-CO- R^{21} , OR^{21} , where

 R^{21} and R^{22} -independently of one another are is hydrogen or C_1 - C_4 -alkyl and

R²³— is hydrogen, C1 C4 alkyl or phenyl

 R^3 is

$$-N$$
 R^{32} , $-N$
 R^{32} , $-N$
 R^{52}

and

 R^{32} is hydrogen and or -(CH₂)_o-(CHR³¹)_m-(CH₂)_n-G, where R^{31} is hydrogen, C₁-C₄-alkyl, OH and or O-C₁-C₄-alkyl,

 m_7 and o independently of one another are m_1 0, 1 or 2 and

n is 1, 2, 3 or 4, and

 R^4 is hydrogen, branched and unbranched C_1 - C_6 -alkyl, chlorine, bromine, fluorine, nitro, cyano, $NR^{41}R^{42}$, NH-CO- R^{43} , or OR^{41} , where

 R^{41} and R^{42} independently of one another are hydrogen or C_1 - C_4 -alkyl and

 R^{43} is C_1 - C_4 -alkyl or phenyl, and

G is NR⁵¹R⁵² or one of the radicals below

where

R⁵¹ is hydrogen and or branched and unbranched C₁-C₆-alkyl and

is hydrogen, COCH₃, CO-O-C₁-C₄-alkyl, COCF₃, or branched and unbranched C₁-C₆-alkyl, it being possible for one hydrogen of the C₁-C₆-alkyl radical to be substituted by one of the following radicals: OH, O-C₁-C₄-alkyl and phenyl and for the phenyl ring also to carry one or two of the following radicals: chlorine, bromine, fluorine, branched and unbranched C₁-C₄-alkyl, nitro, amino, C₁-C₄-alkylamino, OH, O-C₁-C₄-alkyl, and CN, SO₂-C₁-C₄-alkyl,

and the tautomeric forms, a possible enantiomeric and diastereomeric forms thereof, a the prodrugs thereof, or and pharmacologically physiologically tolerated salts thereof.

- 4. (Previously Presented) A compound as claimed in claim 1, where R^2 is in position 3 and R^3 is in position 4 or R^2 is in position 4 and R^3 is in position 3 relative to the benzimidazole ring.
- 5. (Previously Presented) A compound as claimed in claim 1, where R^1 and R^4 are hydrogen.

- 6. (Previously Presented) A compound as claimed in claim 1, where R^2 is hydrogen, branched or unbranched C_1 - C_6 -alkyl, nitro, CN, NH₂, or O- C_1 - C_4 -alkyl.
- 7. (Currently Amended) A compound as claimed in claim 1 where (i) for \mathbb{R}^3 being

 $\mathbb{R}^{34} \mathbb{R}^{32}$ is hydrogen or $-(CH_2)_w - G - (CH_2)_p - G$, where

wp is 1 or 2 and

(ii) for R³ being

 $\mathbb{R}^{31} \mathbb{R}^{32}$ is hydrogen or -(CH₂)_p-G, where

p is 1 or 2 and

and (iii) for R³ being

$$--N$$
 $(CH_2)_{1,2}$ $N--R^{52}$

where

 R^{52} is hydrogen, branched and unbranched C_1 - C_6 -alkyl, where one hydrogen of the C_1 - C_6 -alkyl radical may be substituted by one of the following radicals: OH, O- C_1 - C_4 -alkyl and phenyl, and where the phenyl ring may also carry one or two of

the following radicals: chlorine, bromine, fluorine, branched and unbranched C₁-C₄-alkyl, nitro, amino, C₁-C₄-alkylamino, C₁-C₄-dialkylamino, OH, O-C₁-C₄-alkyl, CN, and SO₂-C₁-C₄-alkyl.

- 8. (Currently Amended) A compound as claimed in claim 1, where R^3 is $-D-(F^1)_p-(E)_q-(F^2)_r-G$ where D is O, F1 F^1 is a C_1-C_4 carbon chain, p is 1, q is 0 and r is 0.
- 9. (Previously Presented) A compound as claimed in claim 1, where R⁵ is a 6-membered ring and R⁵² is an optionally substituted phenyl ring.
- 10. (Previously Presented) A drug comprising besides conventional vehicles and ancillary substances a compound as claimed in claim 1.
- 11. (Previously Presented) A method for treating a disorder in which pathologically elevated PARP activities occur, said method comprising administering an effective amount of a compound of the formula I as claimed in claim 1 to a mammal suffering from said disorder.
- 12. (Currently Amended) The use of compounds of the formula I The method as claimed in claim 11 wherein the disorder is a neurodegenerative disease or involves neuronal damage.
- 13. (Previously Presented) The method as claimed in claim 12, wherein the neurodegenerative disease or neuronal damage is induced by ischemia, trauma or massive

bleeding.

- 14. (Previously Presented) The method as claimed in claim 11 wherein the disorder is stroke or craniocerebral trauma.
- 15. (Currently Amended) The method as claimed in claim 11 wherein the disorder is Alzheimer's disease and or Huntington's disease.

- 16. (Previously Presented) The method as claimed in claim 11 wherein the disorder is damage due to ischemia.
- 17. (Previously Presented) The method as claimed in claim 11 wherein the disorder is epilepsy.
- 18. (Previously Presented) The method as claimed in claim 11 wherein the disorder is damage to the kidneys after renal ischemia, damage caused by drug therapy or damage resulting after kidney transplants.
- 19. (Previously Presented) The method as claimed in claim 11 wherein the disorder is damage to the heart after cardiac ischemia.

- 20. (Currently Amended) The method as claimed in claim 11 wherein the disorder is a microinfarcts microinfarct.
- 21. (Previously Presented) The method as claimed in claim 11 wherein the disorder is under vascularization of critically narrowed coronary arteries.
- 22. (Currently Amended) The method as claimed in claim 11 wherein the disorder is an acute myocardial infarct and or damage during an after medical or mechanical lysis thereof.
- 23. (Previously Presented) The method as claimed in claim 11 wherein the disorder is a tumor or metastasis thereof.
- 24. (Previously Presented) The method as claimed in claim 11 wherein the disorder is sepsis of multi-organ failure.
- 25. (Previously Presented) The method as claimed in claim 11 wherein the disorder is an immunological disease.
- 26. (Previously Presented) The method as claimed in claim 11 wherein the disorder is diabetes mellitus.